

AperTO - Archivio Istituzionale Open Access dell'Università di Torino

Do left atrial and appendage morphology and function help predict thromboembolic risk in atrial fibrillation?

This is the author's manuscript

Original Citation:

Availability:

This version is available <http://hdl.handle.net/2318/1567892> since 2017-09-25T15:24:38Z

Published version:

DOI:10.2459/JCM.0000000000000305

Terms of use:

Open Access

Anyone can freely access the full text of works made available as "Open Access". Works made available under a Creative Commons license can be used according to the terms and conditions of said license. Use of all other works requires consent of the right holder (author or publisher) if not exempted from copyright protection by the applicable law.

(Article begins on next page)



UNIVERSITÀ DEGLI STUDI DI TORINO

This is an author version of the contribution published on:

Questa è la versione dell'autore dell'opera:

Journal of cardiovascular medicine, 17(3), 2016, doi:

10.2459/JCM.0000000000000305]

The definitive version is available at:

La versione definitiva è disponibile alla URL:

<http://journals.lww.com/jcardiovascularmedicine/pages/articleviewer.aspx?year=2016&issue=03000&article=00001&type=abstract>

Do left atrial appendage morphology and function help predict thromboembolic risk in atrial fibrillation??

Matteo Anselmino MD PhD, Sebastiano Gili MD, Davide Castagno MD,

Federico Ferraris MD, Mario Matta MD, Chiara Rovera MD, Carla Giustetto MD, Fiorenzo Gaita

MD Professor

Cardiology Division, Department of Medical Sciences,

University of Turin, Italy

Word count: 2876 words, 3 Figures, 1 Table, 93 references

Key-words: atrial fibrillation; cardiovascular imaging; echocardiography; magnetic resonance;
thromboembolic risk.

Running title: cardiovascular imaging and thromboembolic risk

Corresponding author:

Fiorenzo Gaita, MD Professor

Cardiology Division, Department of Medical Sciences,

University of Turin, Italy

Corso Dogliotti 14, 10126 Turin, Italy

Phone: +39-011-6709557 Fax: +39-011-2369557

Email: fiorenzo.gaita@unito.it

Abstract

Clinical scores (i.e. CHA₂DS₂-VASc) are the mainstay of thromboembolic risk management in nonvalvular atrial fibrillation (AF). Nonetheless, they bear some limitations to precisely define risk-benefit ratio of oral anticoagulation (OAC), both with vitamin K antagonists (VKA) and with novel direct oral anticoagulants (DOAC), especially in patients with low-intermediate scores.

Cardiovascular imaging, allowing to directly visualize those patho-physiological alterations which may lead to the formation of intracardiac thrombi, offers itself as a unique tool helping to refine thromboembolic risk stratification. Many parameters have been tested, focusing primarily on functional and morphological variables of the left atrium (LA) and left atrial appendage (LAA). LA volume and LAA peak flow velocity have, since long-time, been associated with increased thromboembolic risk, while some new parameters, as LA fibrosis assessed by late-gadolinium enhanced (LGE) magnetic resonance imaging (MRI), LA and LAA strain and LAA morphology have more recently shown some ability in predicting embolic events in AF patients. Overall, however, these parameters have seen, to date, a scarce clinical implementation, especially due to inconsistency of validated cut-offs and/or strong clinical evidences driven by technical limitations, such as expensiveness of the technologies (i.e. MRI or computed tomography), invasiveness (i.e. transesophageal echocardiography) or limited reproducibility (i.e. LGE MRI). In conclusion, to date, cardiovascular imaging plays a limited role, however, validation and diffusion of the new techniques hereby systematically presented hold the potential to refine thromboembolic risk stratification in nonvalvular AF.

1 GLOSSARY

2

3D three-dimensional

AF atrial fibrillation

AP antero-posterior

AUC area under the curve

CT computed tomography

DOAC direct oral anticoagulant

LA left atrium

LAA left atrial appendage

LAA_{max} area maximal area of LAA

LAA_v LAA peak flow velocities

LAVI left atrial volume index

LGE late gadolinium enhancement

LVEF *left ventricular ejection fraction*

MRI magnetic resonance imaging

OAC oral anticoagulation

SEC spontaneous echo-contrast

SR sinus rhythm

TDI tissue-Doppler imaging

TEE trans-esophageal echocardiography

TIA transient ischemic attack

TTE trans-thoracic echocardiography

VKA vitamin K antagonist

3

4

1 **Introduction**

2 Atrial fibrillation (AF) is a known risk factor for thromboembolic events, especially stroke and
3 transient ischemic attack (TIA)¹. Indeed, recent data suggest that eliminating the arrhythmia may
4 achieve superior results, compared to simple rate control, in terms of survival², stroke incidence³
5 and decline in cognitive functions⁴. However, when rhythm control strategy is not indicated or fails,
6 oral anticoagulation (OAC), both with vitamin K antagonists (VKA) or novel direct oral
7 anticoagulants (DOAC), is , the only treatment to prevent thromboembolic events. For these reasons
8 on a daily basis, physicians are required to perform an accurate assessment of the risk-benefit ratio
9 for VKAs or each different DOACs⁵ in patients with AF, balancing the implicit bleeding risk.
10 Several clinical scores, such as the CHA₂DS₂-VASc score, have been proposed and more less
11 reliably identify patients potentially benefiting from OAC⁶, considering clinical variables that most
12 strongly predict not only strokes/TIAs but also AF recurrences following transcatheter ablation⁷ and
13 electrical or pharmacological cardioversions⁸. Even if usually reliable, CHA₂DS₂-VASc score still
14 remains inadequate in several clinical circumstances, particularly in patients at low/intermediate
15 risk (i.e., CHA₂DS₂-VASc 1), for whom net clinical benefit of VKAs and even of DOACs is
16 seriously counterbalanced by the bleeding risk^{9,10,11,12}. Trying to overcome this gap in evidence and
17 recommendations, a number of alternative parameters have been investigated, evaluating their value
18 to guide clinical decision-making. Special focus, in fact, has been directed towards the left atrium
19 (LA) and left atrial appendage (LAA), as they harbour the vast majority of thrombi forming during
20 nonvalvular AF^{13,14}.

21 Aim of the present article is to methodologically review these alternative parameters and their
22 application to nonvalvular AF, discussing if and which may aid the physician in everyday activity.

23

24

25

1 **Spontaneous echocontrast and thrombi**

2 Any discussion concerning LA and LAA and the risk of thromboembolic events in patients with AF
3 has to deal first with the two strongest predictors of cardioembolic stroke, the presence of thrombi
4 or spontaneous echo-contrast (SEC) in the LA or LAA¹⁵. The relationship between these findings
5 and the risk of subsequent cardioembolic stroke is known since the dawning of echocardiography.
6 Presence of thrombi represents the ultimate stage before the clinical event, even if their presence
7 doesn't necessarily end in peripheral embolization^{16,17}. SEC represents instead the most obvious
8 visual representation of the blood stasis occurring during atrial dysrhythmias. Its presence is graded,
9 depending from intensity, on a scale ranging from 0 to 4 (0, absence of SEC; 4, severe SEC with
10 intense echodensity and very slow swirling patterns in the LAA and usually LA; 1, 2 and 3 mild,
11 mild to moderate and moderate SEC, respectively, with intermediate features compared to 0 and
12 4)¹⁸, and its relationship with subsequent embolic events has been consistently demonstrated^{19,20}.

13 Prevalence of thrombi is highly variable depending on clinical setting: in consecutive,
14 anticoagulated patients undergoing AF ablation, they have been reported in 0.6%-3.6% of the
15 patients^{21,22}, while higher figures have been described in different clinical settings (up to 5.9%-
16 16.5%^{23,24,25}). Prevalence of SEC, also varies widely: for example it ranged from 35% in the large
17 cohort of anticoagulated patients undergoing AF ablation enrolled by Puwanant et al. (80% of
18 patients with CHADS₂ score 0-1, 13% valvular AF, 9% with left ventricular ejection fraction
19 [LVEF] < 35%)²¹, to 8% in the study by Kleemann and colleagues (nonvalvular AF, CHADS₂ score
20 0-1).

21 Despite their strong and consistent ability to predict stroke and other cardioembolic events, these
22 parameters, however, present, at least, two main limitations: first, they represent an advanced grade
23 of prothrombotic state, the last step before a clinical event, so that they're rarely useful for patients
24 with recent onset of AF or at low/intermediate clinical risk in whom a decision on OAC

prescription has to be made; second, these alterations are recorded by trans-esophageal echocardiography (TEE) and only rarely by trans-thoracic echocardiography (TTE). For these reasons, other parameters have been assessed, often referring to thrombi or SEC as surrogate study end-points.

Morphological and structural assessment

Left atrium

During AF, LA enlargement is a common finding, as it comes both as a cause and as a consequence of the arrhythmia^{26,27}. Since echocardiography became available, a correlation between LA dimensions as measured by antero-posterior (AP) diameter and thromboembolic risk was sought, resulting in contradictory data^{28,29,30,31}. In 1995, data from a prospective cohort of 1,371 men and 1,728 women (of whom approximately 1.8% with AF) included in the Framingham study reported a significant association between LA AP diameter as measured by M-mode echocardiography and the risk of stroke and death. The relative risk of stroke was 2.4 per 10 mm increment in men (95% CI, 1.6 to 3.7) and 1.4 in women (95% CI, 0.9 to 2.1), and, when controlling for prevalent and interim AF, it remained significant, even if with a weaker association (2.0 for men, 1.2 for women)³². More recently a study on 500 consecutive patients, admitted to hospital for stroke or TIA (15.6% with AF), confirmed LA AP diameter as an independent predictor for SEC/thrombi³³ and LA AP diameter, evaluated by multi-detector computed tomography (CT) in 67 patients admitted to hospital for stroke was found to be significantly increased as compared to AF controls without stroke³⁴. On the other side, a meta-analysis performed on individual data from 1,066 patients enrolled in three large clinical trials from late 80s ó early 90s^{35,36,37} showed that LA AP diameter did not predict stroke neither at univariate analysis nor when adjusted for clinical predictors³⁸. Similar results arose from a prospective study assessing the prognostic implications of SEC¹⁵ and by a post-hoc analysis³⁹ of the Atrial Fibrillation Follow-up Investigation of Rhythm Management

1 (AFFIRM) trial⁴⁰, a study in which 4,060 patients with recurrent AF were randomised to rhythm
2 control strategy vs. rate control strategy.

3 Given these data inconsistencies it becomes evident that other LA parameters need to be assessed in
4 relationship to thromboembolic events. LA area related, in AF patients, to TEE high-risk features
5 for thromboembolic events⁴¹ and, when integrated with LV dysfunction, evidently enhanced the
6 performance of CHADS₂ and CHA₂DS₂-VASc clinical scores to recognize patients at high
7 thromboembolic risk⁴². However, also for LA area data are contradictory, as, for example, in the
8 1994 study by Fatkin and colleagues, no significant relationship arose between LA area and SEC or
9 LA thrombi¹⁸.

10 LA volume assessed by transthoracic echocardiography has been more widely tested, and,
11 especially when indexed for body surface (left atrial volume index, LAVI), it showed more
12 consistent interaction with clinical events. LA volume, both total and indexed for body surface,
13 significantly related to LAA thrombi at TEE examination^{41,43,44}. In two distinct studies from
14 populations based in Olmsted County, increased LAVI ($\times 32$ ml/mq) related to cardiovascular
15 adverse events: in the first report, enrolling 1,160 elderly patients (75.5 years) in sinus rhythm (SR),
16 the predictive value of LAVI was validated in a multivariate model⁴⁵, while in the second study,
17 which involved 46 young adults (< 60 years) with lone AF followed for a median of 27 years, the
18 association was confirmed only in a univariate setting, due to the low number of index events⁴⁶.

19 Although suggestive LA enlargement is strictly dependent on its underlying cause. Indeed, Ayrala
20 and colleagues reported that the predictive value of LAVI for LAA thrombi at TEE was stronger
21 when LA enlargement was associated with a reduction in LVEF⁴⁷, while, in an Olmsted County-
22 based cohort of 2,042 patients with more than 45 years (3% with a history of AF), Pritchett et al.
23 found that the relationship between LAVI and mortality did not persist when controlling for
24 diastolic dysfunction⁴⁸, a possible predictor *per se* of embolic events in AF⁴⁹. Despite the

1 aforementioned data, however, to date, no conclusive evidence supports the use of LA enlargement
2 markers to predict embolic events in AF.

3 Eventually, progresses in magnetic resonance imaging (MRI), particularly with the development of
4 late-gadolinium enhancement (LGE) imaging, have been employed to assess structural changes of
5 atrial myocardium. Particularly, a dedicated protocol, first developed by a research group based in
6 Salt Lake City, allowed to semi-quantify LA fibrosis and to classify it in four classes based on its
7 extension (Utah classes I-IV)⁵⁰. LA LGE extent assessed by this method related to AF recurrence
8 after ablation⁵¹ and, more interestingly, to the presence of thrombi detected by TEE⁵² and with a
9 medical history of previous stroke⁵³ (**figure 1**). Despite its brilliant promises, this technique has
10 embraced to date little diffusion, probably due to the high costs of MRI and the absence of
11 prospective validation⁵⁴. This type of analysis is moreover penalised by the development of
12 different algorithms in different centres, a substantial limit to the standardisation of the results. For
13 this reason, a public challenge was launched trying to determine the better performing algorithm
14 and, even if no definite results were reached, this attempt may be a foundation stone in the diffusion
15 and widespread adoption of this evaluation⁵⁵.

16 17 *Left atrial appendage*

18 Concerning LAA dimensions, interpretation of existing literature is, unfortunately, even more
19 inconclusive. A paucity of data has been published and, since LAA measurements are far less
20 standardized than LA, generalizations of results and comparisons are indeed challenging.

21 Usually, LAA abnormalities are related to LA alterations^{56,57,58}, even if some peculiar traits can be
22 found, making a dedicated analysis of LAA worth a chance. Indeed, LAA enlargement doesn't
23 always coincide with increased LA size (and viceversa)⁵⁹, and, at least in patients in SR, LAA peak

1 flow velocity doesn't relate to any echocardiographic parameter connected with LA except than Aø
2 measured at tissue-Doppler imaging (TDI)⁶⁰.

3 Pollick and colleagues reported that a bigger maximal LAA (LAA_{max}) area (and not LA AP
4 diameter) as measured by TEE related to the presence of thrombi and/or SEC in 19 non-
5 anticoagulated patients with valvular and nonvalvular AF⁵⁹, and, while another study confirmed this
6 finding⁶¹, other laboratories failed to consistently relate LAA area to an increased risk of stroke⁶² or
7 thromboembolic events^{63,64}.

8 LAA size analysis by CT or MRI also generally did not provide strong evidences in favour of a role
9 of LAA volume. Two studies reported a direct correlation between increased LAA volumes and
10 previous stroke, even after controlling for possible confounding factors, with an ideal cut-off for
11 significant risk increase at 34 ml³ ⁶⁵, but the majority of experiences did not confirm this
12 relationship^{66,67,68,69}.

13 More recently, LAA morphology, by CT or MRI, has been proposed to be described in four
14 different morphologies (Cactus, Chicken Wing, Windsock and Cauliflower) based on the
15 characteristics of the main and secondary lobes⁷⁰. By this categorization LAA morphology related
16 to thromboembolic events: it was first demonstrated that, in patients undergoing AF transcatheter
17 ablation, Chicken Wing, the most frequent LAA morphology, independently related to the absence
18 of previous stroke/TIA⁶⁹, and that the prevalence of silent cerebral ischemic lesions increased with
19 growing complexity of LAA morphology, from Cactus, the more simple, through Chicken Wing,
20 Windsock and Cauliflower, the more complex⁷¹ (**figure 2**). Once again, further studies brought
21 controversial results: two experiences, indeed, confirmed the existence of a relationship between
22 LAA morphology and cerebrovascular events, even if with some slight differences (Kimura and
23 colleagues identified Cauliflower as the high-risk morphology⁷², Kosiuk et al. Chicken Wing⁷³),
24 while other studies failed to report any correlation between LAA morphology and previous stroke⁷⁴

and incident stroke or TIA after AF ablation⁶⁷. Concerning LAA morphology the main limitations seems to relate to the reproducibility of the classification by CT or MRI⁷⁴. More interestingly, however, a recent study by Petersen and colleagues, assessed LAA morphology by three-dimensional (3D) TEE, which resulted non-inferior to CT and MRI [*data in press*], and found a correlation, corroborated by multivariate linear regression analysis, between non-Chicken Wing LAA morphologies and reduced LAA flow velocities⁷⁵. These latter studies are particularly promising, as validation of LAA morphology description by echocardiographic techniques may enhance the routine clinical implementation of this analysis, helping to refine thromboembolic risk stratification.

Functional assessment

Left atrium

LA and LAA function impairment that lead to blood stasis are, indeed, at the base of thrombus formation⁷⁶.

As for LA size, an impaired LA function has been associated with an unfavourable outcome in non-AF patients: a reduced LA function index (a multiparametric index incorporating LA size, LA emptying fraction and LV outflow velocity-time integral) significantly related to incident stroke or TIA at follow up in 893 patients without a known history of AF not on OAC⁷⁷, while a reduced LA longitudinal strain, as assessed by speckle tracking, was able to predict development of adverse events including, among others, AF, stroke or TIA and cardiovascular death in 312 patients in SR⁷⁸. More in details, within AF patients, CHADS₂ and CHA₂DS₂-VASc scores significantly related to an increased LAVI and a reduced LA emptying fraction (LAEF) in a sub-analysis of the ENGAGE AF-TIMI 48 study. LAVI and LAEF also inversely related to each other, even if only LAVI was confirmed as an independent predictor after adjusting for confounding factors⁷⁹. An association between LA function and CHADS₂ and CHA₂DS₂-VASc scores was also confirmed when LAEF

1 was assessed by 3D echocardiography⁸⁰.

2 In addition, since its introduction and widespread diffusion, strain analysis by speckle tracking has

3 been increasingly applied also in this field, reporting very promising results. Even if a prospective

4 validation has yet to be provided, a reduced global longitudinal strain proved to relate to the

5 extent of fibrosis at MRI⁸¹ and not only to an increased CHADS₂ or CHA₂DS₂-VASc score^{80,82,83},

6 but to an improved CHADS₂ and CHA₂DS₂-VAScscores ability to identify patients with recent

7 acute embolic events and to predict mortality at follow up^{83,84}. Interestingly, peak negative and

8 positive atrial strain were able to discriminate patients with previous stroke or TIA in an AF-cohort

9 with low intermediate CHADS₂ score (Ö1, of whom 30% with CHA₂DS₂-VAScÖ1)⁸⁵.

10

11 *Left atrial appendage*

12 LAA function has been thoroughly investigated since a correlation between reduced LAA peak

13 flow velocities (LAAv) and increased thromboembolic risk was demonstrated. LAAv tend to reduce

14 during the course of AF and more decreased values strongly related to the presence of SEC¹⁸, and

15 of LAA thrombi³³ and predicted cardioembolic events in a sub-analysis of the SPAF-III study⁸⁶.

16 Despite overall consistent results, this parameter has embraced a very limited clinical

17 implementation, partly due to the variable and widely ranging cut-offs reported (in the 3

18 aforementioned studies, LAAv relating to adverse events were < 35 cm/s, < 55 cm/s and < 20 cm/s,

19 respectively^{18,33,86}), and, partly, because LAAv measurement require TEE.

20 To overcome this latter limitation, surrogate markers of reduced LAAv have been investigated. As

21 TEE studies reported a correlation between reduced LAA wall acceleration as measured by TDI and

22 increased thromboembolic risk^{87,88}, an evaluation of those TDI parameters by TTE has been

23 attempted, with garbed success. With a good feasibility (> 90%), LAA wall velocity, evaluated

24 mainly by LAA tip acceleration on a short-axis view, significantly related to reduced LAAv,

25 presence of SEC or thrombi and previous stroke^{89,90,91} (**figure 3**). Moreover, these results were

26 prospectively validate in a cohort of 179 patients with previous thromboembolic stroke related to

AF, in whom an LAA wall velocity < 8.7 cm/s independently predicted recurrent stroke or cerebrovascular death⁹². TDI was also used to assess septal and lateral mitral annular aø wave velocities, which strongly related to reduced LAAv and SEC^{60,93}. Eventually a similar proceeding than with TDI was attempted with LAA strain, even if data are, to date, scarce. In fact, LAA strain predicted the presence of LAA thrombi²⁵, and feasibility of the evaluation by TTE was confirmed in a series of 82 patients, in whom negative peak strain rate and time-to-peak positive strain independently related to the presence of LAA thrombi or sludge with an AUC of 0.89⁹⁴. Finally, LAA emptying determined by velocity encoded-MRI had a good correlation with LAAv as measured by Pulsed Doppler at TEE, opening the doors of LAA functional evaluation to a new technique⁹⁵.

Conclusion

LAVI and functional parameters of both LA and LAA consistently predicted stroke in patients with AF, while promising results came from the application of more recent techniques such as strain and LAA morphology assessment.

Clinical application of these parameters, however, is to date limited, mostly due to lack of large-scale prospective validation and consequent inability to suggest significant cut-offs.

In our opinion, to date, parameters, such as severely reduced LAAv or markedly enlarged LA, may aid, in a case-by-case approach, as complementary tools to guide OAC prescription in controversial and ambiguous cases. In addition, LAA and LA imaging, particularly with the development of technologies such as TDI or speckle tracking, bears a promising potential.

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21

Conflicts of interest: no conflicts of interest pertaining the present work have been reported.

1 References

-
- ¹ Wolf PA, Dawber TR, Thomas HE Jr, Kannel WB. Epidemiologic assessment of chronic atrial fibrillation and risk of stroke: the Framingham study. *Neurology*. 1978;28(10):973-7
- ² Ionescu-Ittu R, Abrahamowicz M, Jackevicius CA, et al. Comparative effectiveness of rhythm control vs rate control drug treatment effect on mortality in patients with atrial fibrillation. *Arch Intern Med* 2012;**172**(13):997-1004
- ³ Tsadok MA, Jackevicius CA, Essebag V, et al. Rhythm versus rate control therapy and subsequent stroke or transient ischemic attack in patients with atrial fibrillation. *Circulation* 2012;126(23):2680-7
- ⁴ Gaita F, Corsinovi L, Anselmino M, et al. Prevalence of silent cerebral ischemia in paroxysmal and persistent atrial fibrillation and correlation with cognitive function. *J Am Coll Cardiol* 2013;62(21):1990-7
- ⁵ Prisco D, Cenci C, Silvestri E, Ciucciarelli L, Di Minno G. Novel oral anticoagulants in atrial fibrillation: which novel oral anticoagulant for which patient? *J Cardiovasc Med (Hagerstown)*. 2015 Mar 19. [Epub ahead of print]
- ⁶ Lip GY, Nieuwlaat R, Pisters R, Lane DA, Crijns HJ. Refining clinical risk stratification for predicting stroke and thromboembolism in atrial fibrillation using a novel risk factor-based approach: the euro heart survey on atrial fibrillation. *Chest*. 2010;137(2):263-72
- ⁷ Jacobs V, May HT, Bair TL, et al. The impact of risk score (CHADS2 versus CHA2DS2-VASc) on long-term outcomes after atrial fibrillation ablation. *Heart Rhythm*. 2015;12(4):681-6
- ⁸ Falsetti L, Viticchi G, Tarquinio N, Silvestrini M, Capeci W, Balloni A, Catozzo V, Gentile A, Pellegrini F. CHA2DS2-VASc in the prediction of early atrial fibrillation relapses after electrical or pharmacological cardioversion. *J Cardiovasc Med (Hagerstown)*. 2014;15(8):636-41
- ⁹ Chao TF, Liu CJ, Wang KL, et al. Should Atrial Fibrillation Patients With 1 Additional Risk Factor of the CHA2DS2-VASc Score (Beyond Sex) Receive Oral Anticoagulation? *J Am Coll*

Cardiol. 2015;65(7):635-42

¹⁰ Banerjee A, Lane DA, Torp-Pedersen C, Lip GY. Net clinical benefit of new oral anticoagulants (dabigatran, rivaroxaban, apixaban) versus no treatment in a 'real world' atrial fibrillation population: a modelling analysis based on a nationwide cohort study. *Thromb Haemost.* 2012;107(3):584-9

¹¹ Friberg L, Skeppholm M, Terént A. Benefit of Anticoagulation Unlikely in Patients With Atrial Fibrillation and a CHA2DS2-VASc Score of 1. *J Am Coll Cardiol.* 2015;65(3):225-32

¹² Calkins H. Data Strengthen to Support Recommending Anticoagulant Therapy for All Atrial Fibrillation Patients With a CHA2DS2-VASc Score ≥ 1 . *J Am Coll Cardiol.* 2015;65(7):643-4

¹³ Blackshear JL, Odell JA. Appendage obliteration to reduce stroke in cardiac surgical patients with atrial fibrillation. *Ann Thorac Surg.* 1996;61(2):755-9

¹⁴ Bogousslavsky J, Van Melle G, Regli F, Kappenberger L. Pathogenesis of anterior circulation stroke in patients with nonvalvular atrial fibrillation: the Lausanne Stroke Registry. *Neurology.* 1990;40(7):1046-50

¹⁵ Leung DY, Black IW, Cranney GB, Hopkins AP, Walsh WF. Prognostic implications of left atrial spontaneous echo contrast in nonvalvular atrial fibrillation. *J Am Coll Cardiol.* 1994;24(3):755-62

¹⁶ Stöhlberger C, Chnupa P, Kronik G, et al. Transesophageal echocardiography to assess embolic risk in patients with atrial fibrillation. ELAT Study Group. Embolism in Left Atrial Thrombi. *Ann Intern Med.* 1998;128(8):630-8

¹⁷ Stoddard MF, Singh P, Dawn B, Longaker RA. Left atrial thrombus predicts transient ischemic attack in patients with atrial fibrillation. *Am Heart J.* 2003;145(4):676-82

¹⁸ Fatkin D, Kelly RP, Feneley MP. Relations between left atrial appendage blood flow velocity, spontaneous echocardiographic contrast and thromboembolic risk in vivo. *J Am Coll Cardiol.* 1994;23(4):961-9

-
- ¹⁹ Transesophageal echocardiographic correlates of thromboembolism in high-risk patients with nonvalvular atrial fibrillation. The Stroke Prevention in Atrial Fibrillation Investigators Committee on Echocardiography. *Ann Intern Med.* 1998;128(8):639-47
- ²⁰ Lowe BS, Kusunose K, Motoki H, et al. Prognostic significance of left atrial appendage "sludge" in patients with atrial fibrillation: a new transesophageal echocardiographic thromboembolic risk factor. *J Am Soc Echocardiogr.* 2014;27(11):1176-83
- ²¹ Puwanant S, Varr BC, Shrestha K, et al. Role of the CHADS2 score in the evaluation of thromboembolic risk in patients with atrial fibrillation undergoing transesophageal echocardiography before pulmonary vein isolation. *J Am Coll Cardiol* 2009;54(22):2032-9
- ²² Wallace TW, Atwater BD, Daubert JP, et al. Prevalence and clinical characteristics associated with left atrial appendage thrombus in fully anticoagulated patients undergoing catheter-directed atrial fibrillation ablation. *J Cardiovasc Electrophysiol* 2010;21(8):849-52
- ²³ Maltagliati A, Galli CA, Tamborini G, Celeste F, Muratori M, Pepi M. Incidence of spontaneous echocontrast, 'sludge' and thrombi before cardioversion in patients with atrial fibrillation: new insights into the role of transesophageal echocardiography. *J Cardiovasc Med (Hagerstown).* 2009;10(7):523-8
- ²⁴ Seidl K, Rameken M, Drögemüller A, et al. Embolic events in patients with atrial fibrillation and effective anticoagulation: value of transesophageal echocardiography to guide direct-current cardioversion. Final results of the Ludwigshafen Observational Cardioversion Study. *J Am Coll Cardiol.* 2002;39(9):1436-42
- ²⁵ Ono K, Iwama M, Kawasaki M, et al. Motion of left atrial appendage as a determinant of thrombus formation in patients with a low CHADS2 score receiving warfarin for persistent nonvalvular atrial fibrillation. *Cardiovasc Ultrasound.* 2012;10:50.

-
- ²⁶ Howard C, Dittrich, MD, Lesly A, et al. on behalf of the Stroke Prevention in Atrial Fibrillation Investigators. Left atrial diameter in nonvalvular atrial fibrillation: An echocardiographic study. *Am Heart J* 1999;137:494-9
- ²⁷ Psaty BM, Manolio TA, Kuller LH, et al. Incidence of and risk factors for atrial fibrillation in older adults. *Circulation*. 1997;96(7):2455-61
- ²⁸ Caplan LR, D'Cruz I, Hier DB, Reddy H, Shah S. Atrial size, atrial fibrillation, and stroke. *Ann Neurol*. 1986;19(2):158-61.
- ²⁹ Petersen P, Kastrup J, Helweg-Larsen S, Boysen G, Godtfredsen J. Risk factors for thromboembolic complications in chronic atrial fibrillation. The Copenhagen AFASAK study. *Arch Intern Med*. 1990;150(4):819-21
- ³⁰ Gustafsson C, Britton M, Brolund F, Eriksson SV, Lindvall K. Echocardiographic findings and the increased risk of stroke in nonvalvular atrial fibrillation. *Cardiology*. 1992;81(4-5):189-95
- ³¹ Corbalán R, Arriagada D, Braun S, et al. Risk factors for systemic embolism in patients with paroxysmal atrial fibrillation. *Am Heart J*. 1992;124(1):149-53
- ³² Benjamin EJ, D'Agostino RB, Belanger AJ, Wolf PA, Levy D. Left atrial size and the risk of stroke and death. The Framingham Heart Study. *Circulation*. 1995;92(4):835-41.
- ³³ Handke M, Harloff A, Hetzel A, Olschewski M, Bode C, Geibel A. Left atrial appendage flow velocity as a quantitative surrogate parameter for thromboembolic risk: determinants and relationship to spontaneous echocontrast and thrombus formation--a transesophageal echocardiographic study in 500 patients with cerebral ischemia. *J Am Soc Echocardiogr*. 2005;18(12):1366-72
- ³⁴ Lee JM, Shim J, Uhm JS, et al. Impact of increased orifice size and decreased flow velocity of left atrial appendage on stroke in nonvalvular atrial fibrillation. *Am J Cardiol*. 2014;113(6):963-9
- ³⁵ Stroke Prevention in Atrial Fibrillation Study. Final results. *Circulation* 1991;84(2):527-39.

-
- ³⁶ Ezekowitz MD, Bridgers SL, James KE, et al. Warfarin in the prevention of stroke associated with nonrheumatic atrial fibrillation. Veterans Affairs Stroke Prevention in Nonrheumatic Atrial Fibrillation Investigators. *N Engl J Med* 1992;327(20):1406-12.
- ³⁷ The effect of low-dose warfarin on the risk of stroke in patients with nonrheumatic atrial fibrillation. The Boston Area Anticoagulation Trial for Atrial Fibrillation Investigators. *N Engl J Med* 1990;323(22):1505-11
- ³⁸ Echocardiographic predictors of stroke in patients with atrial fibrillation: a prospective study of 1066 patients from 3 clinical trials. *Arch Intern Med*. 1998;158(12):1316-20
- ³⁹ Olshansky B, Heller EN, Mitchell LB, et al. Are transthoracic echocardiographic parameters associated with atrial fibrillation recurrence or stroke? Results from the Atrial Fibrillation Follow-Up Investigation of Rhythm Management (AFFIRM) study. *J Am Coll Cardiol*. 2005;45(12):2026-33
- ⁴⁰ Wyse DG, Waldo AL, DiMarco JP, et al. Atrial Fibrillation Follow-up Investigation of Rhythm Management (AFFIRM) Investigators. A comparison of rate control and rhythm control in patients with atrial fibrillation. *N Engl J Med*. 2002;347(23):1825-33
- ⁴¹ Faustino A, Providência R, Barra S, et al. Which method of left atrium size quantification is the most accurate to recognize thromboembolic risk in patients with non-valvular atrial fibrillation? *Cardiovasc Ultrasound*. 2014;12:28
- ⁴² Providência R, Botelho A, Trigo J, et al. Possible refinement of clinical thromboembolism assessment in patients with atrial fibrillation using echocardiographic parameters. *Europace* 2012;14(1):36-45
- ⁴³ Wysokinski WE, Ammash N, Sobande F, Kalsi H, Hodge D, McBane RD. Predicting left atrial thrombi in atrial fibrillation. *Am Heart J*. 2010;159(4):665-71
- ⁴⁴ Yamamoto M, Seo Y, Kawamatsu N, et al. Complex left atrial appendage morphology and left atrial appendage thrombus formation in patients with atrial fibrillation. *Circ Cardiovasc Imaging*.

- ⁴⁵ Tsang TS, Barnes ME, Gersh BJ, et al. Prediction of risk for first age-related cardiovascular events in an elderly population: the incremental value of echocardiography. *J Am Coll Cardiol*. 2003;42(7):1199-205
- ⁴⁶ Osranek M, Bursi F, Bailey KR, et al. Left atrial volume predicts cardiovascular events in patients originally diagnosed with lone atrial fibrillation: three-decade follow-up. *Eur Heart J*. 2005;26(23):2556-61
- ⁴⁷ Ayirala S, Kumar S, O'Sullivan DM, Silverman DI. Echocardiographic predictors of left atrial appendage thrombus formation. *J Am Soc Echocardiogr*. 2011;24(5):499-505
- ⁴⁸ Pritchett AM, Mahoney DW, Jacobsen SJ, Rodeheffer RJ, Karon BL, Redfield MM. Diastolic dysfunction and left atrial volume: a population-based study. *J Am Coll Cardiol*. 2005;45(1):87-92
- ⁴⁹ Lee SH, Choi S, Chung WJ, et al. Tissue Doppler index, E/E', and ischemic stroke in patients with atrial fibrillation and preserved left ventricular ejection fraction. *J Neurol Sci*. 2008;271(1-2):148-52
- ⁵⁰ Akoum N, Daccarett M, McGann C, et al. Atrial fibrosis helps select the appropriate patient and strategy in catheter ablation of atrial fibrillation: a DE-MRI guided approach. *J Cardiovasc Electrophysiol*. 2011;22(1):16-22.
- ⁵¹ Oakes RS, Badger TJ, Kholmovski EG, et al. Detection and quantification of left atrial structural remodeling with delayed-enhancement magnetic resonance imaging in patients with atrial fibrillation. *Circulation*. 2009;119(13):1758-67
- ⁵² Akoum N, Fernandez G, Wilson B, McGann C, Kholmovski E, Marrouche N. Association of atrial fibrosis quantified using LGE-MRI with atrial appendage thrombus and spontaneous contrast on transesophageal echocardiography in patients with atrial fibrillation. *J Cardiovasc Electrophysiol*. 2013;24(10):1104-9.
- ⁵³ Daccarett M, Badger TJ, Akoum N, et al. Association of left atrial fibrosis detected by delayed-

-
- enhancement magnetic resonance imaging and the risk of stroke in patients with atrial fibrillation. *J Am Coll Cardiol*. 2011;57(7):831-8
- ⁵⁴ Gaita F, Blandino A. Atrial fibrillation. Left atrial fibrosis--a promising stroke risk factor? *Nat Rev Cardiol*. 2011;8(6):307-8.
- ⁵⁵ Karim R, Housden RJ, Balasubramaniam M, et al. Evaluation of current algorithms for segmentation of scar tissue from late gadolinium enhancement cardiovascular magnetic resonance of the left atrium: an open-access grand challenge. *J Cardiovasc Magn Reson*. 2013;15:105
- ⁵⁶ Anselmino M, Blandino A, Beninati S, et al. Morphologic analysis of left atrial anatomy by magnetic resonance angiography in patients with atrial fibrillation: a large single center experience. *J Cardiovasc Electrophysiol*. 2011;22(1):1-7
- ⁵⁷ Uslu N, Nurkalem Z, Orhan AL, et al. Transthoracic echocardiographic predictors of the left atrial appendage contraction velocity in stroke patients with sinus rhythm. *Tohoku J Exp Med*. 2006;208(4):291-8
- ⁵⁸ Igarashi Y, Kashimura K, Makiyama Y, Sato T, Ojima K, Aizawa Y. Left atrial appendage dysfunction in chronic nonvalvular atrial fibrillation is significantly associated with an elevated level of brain natriuretic peptide and a prothrombotic state. *Jpn Circ J*. 2001;65(9):788-92
- ⁵⁹ Pollick C, Taylor D. Assessment of left atrial appendage function by transesophageal echocardiography. Implications for the development of thrombus. *Circulation*. 1991;84(1):223-31
- ⁶⁰ Agmon Y, Khandheria BK, Meissner I, et al. Are left atrial appendage flow velocities adequate surrogates of global left atrial function? A population-based transthoracic and transesophageal echocardiographic study. *J Am Soc Echocardiogr*. 2002;15(5):433-40
- ⁶¹ Ozer N, Kiliç H, Arslan U, et al. Echocardiographic predictors of left atrial appendage spontaneous echocontrast in patients with stroke and atrial fibrillation. *J Am Soc Echocardiogr*. 2005;18(12):1362-5
- ⁶² Zabalgoitia M, Halperin JL, Pearce LA, Blackshear JL, Asinger RW, Hart RG. Transesophageal

echocardiographic correlates of clinical risk of thromboembolism in nonvalvular atrial fibrillation.

Stroke Prevention in Atrial Fibrillation III Investigators. *J Am Coll Cardiol.* 1998;31(7):1622-6

⁶³ Miyazaki S, Ito T, Suwa M, Nakamura T, Kobashi A, Kitaura Y. Role of transesophageal echocardiography in the prediction of thromboembolism in patients with chronic nonvalvular atrial fibrillation. *Jpn Circ J.* 2001;65(10):874-8

⁶⁴ Okuyama H, Hirono O, Tamura H, et al. Usefulness of intensity variation in the left atrial appendage with contrast echocardiography to predict ischemic stroke recurrence in patients with atrial fibrillation. *Am J Cardiol.* 2008;101(11):1630-7

⁶⁵ Burrell LD, Horne BD, Anderson JL, Muhlestein JB, Whisenant BK. Usefulness of left atrial appendage volume as a predictor of embolic stroke in patients with atrial fibrillation. *Am J Cardiol.* 2013;112(8):1148-52

⁶⁶ Kosiuk J, Nedios S, Kornej J, et al. Impact of left atrial appendage morphology on peri-interventional thromboembolic risk during catheter ablation of atrial fibrillation. *Heart Rhythm.* 2014;11(9):1522-7

⁶⁷ Nedios S, Kornej J, Koutalas E, et al. Left atrial appendage morphology and thromboembolic risk after catheter ablation for atrial fibrillation. *Heart Rhythm.* 2014;11(12):2239-46

⁶⁸ Beinart R, Heist EK, Newell JB, Holmvang G, Ruskin JN, Mansour M. Left atrial appendage dimensions predict the risk of stroke/TIA in patients with atrial fibrillation. *J Cardiovasc Electrophysiol.* 2011;22(1):10-5

⁶⁹ Di Biase L, Santangeli P, Anselmino M, et al. Does the left atrial appendage morphology correlate with the risk of stroke in patients with atrial fibrillation? Results from a multicenter study. *J Am Coll Cardiol.* 2012;60(6):531-8

⁷⁰ Wang Y, Di Biase L, Horton RP, Nguyen T, Morhanty P, Natale A. Left atrial appendage studied by computed tomography to help planning for appendage closure device placement. *J Cardiovasc Electrophysiol.* 2010;21(9):973-82.

-
- ⁷¹ Anselmino M, Scaglione M, Di Biase L, et al. Left atrial appendage morphology and silent cerebral ischemia in patients with atrial fibrillation. *Heart Rhythm*. 2014;11(1):2-7.
- ⁷² Kimura T, Takatsuki S, Inagawa K, et al. Anatomical characteristics of the left atrial appendage in cardiogenic stroke with low CHADS2 scores. *Heart Rhythm*. 2013;10(6):921-5.
- ⁷³ Kosiuk J, Nedios S, Kornej J, et al. Impact of left atrial appendage morphology on peri-interventional thromboembolic risk during catheter ablation of atrial fibrillation. *Heart Rhythm*. 2014;11(9):1522-7
- ⁷⁴ Khurram IM, Dewire J, Mager M, et al. Relationship between left atrial appendage morphology and stroke in patients with atrial fibrillation. *Heart Rhythm*. 2013;10(12):1843-9
- ⁷⁵ Petersen M, Roehrich A, Balzer J, et al. Left atrial appendage morphology is closely associated with specific echocardiographic flow pattern in patients with atrial fibrillation. *Europace*. 2014. pii: euu347.
- ⁷⁶ Watson T, Shantsila E, Lip GY. Mechanisms of thrombogenesis in atrial fibrillation: Virchow's triad revisited. *Lancet*. 2009;373(9658):155-66.
- ⁷⁷ Wong JM, Welles CC, Azarbal F, Whooley MA, Schiller NB, Turakhia MP. Relation of left atrial dysfunction to ischemic stroke in patients with coronary heart disease (from the heart and soul study). *Am J Cardiol*. 2014;113(10):1679-84
- ⁷⁸ Cameli M, Lisi M, Focardi M, et al. Left atrial deformation analysis by speckle tracking echocardiography for prediction of cardiovascular outcomes. *Am J Cardiol*. 2012;110(2):264-9
- ⁷⁹ Gupta DK, Shah AM, Giugliano RP, et al. Effective anticoagulation with factor xA next Generation in AF-Thrombolysis In Myocardial Infarction 48 Echocardiographic Study Investigators. Left atrial structure and function in atrial fibrillation: ENGAGE AF-TIMI 48. *Eur Heart J*. 2014;35(22):1457-65
- ⁸⁰ Islas F, Olmos C, Vieira C, et al. Thromboembolic Risk in Atrial Fibrillation: Association between Left Atrium Mechanics and Risk Scores. A Study Based on 3D Wall-Motion Tracking

Technology. Echocardiography. 2014. doi: 10.1111/echo.12711.

⁸¹ Kuppahally SS, Akoum N, Burgon NS, et al. Left atrial strain and strain rate in patients with paroxysmal and persistent atrial fibrillation: relationship to left atrial structural remodeling detected by delayed-enhancement MRI. *Circ Cardiovasc Imaging*. 2010;3(3):231-9

⁸² Li Y, Ding W, Wang H, et al. Relationship of CHA₂DS₂-VASc and CHADS₂ Score to Left Atrial Remodeling Detected by Velocity Vector Imaging in Patients with Atrial Fibrillation. *PLoS ONE* 8(10): e77653. doi:10.1371/journal.pone.0077653

⁸³ Saha SK, Anderson PL, Caracciolo G, et al. Global Left Atrial Strain Correlates with CHADS₂ Risk Score in Patients with Atrial Fibrillation. *J Am Soc Echocardiogr* 2011;24:506-12

⁸⁴ Obokata M, Negishi K, Kurosawa K, et al. Left atrial strain provides incremental value for embolism risk stratification over CHA₂DS₂-VASc score and indicates prognostic impact in patients with atrial fibrillation. *J Am Soc Echocardiogr*. 2014;27(7):709-716

⁸⁵ Azemi T, Rabdiya VM, Ayirala SR, McCullough LD, Silverman DI. Left atrial strain is reduced in patients with atrial fibrillation, stroke or TIA, and low risk CHADS(2) scores. *J Am Soc Echocardiogr*. 2012;25(12):1327-32

⁸⁶ Goldman ME, Pearce LA, Hart RG, et al. Pathophysiologic correlates of thromboembolism in nonvalvular atrial fibrillation: I. Reduced flow velocity in the left atrial appendage (The Stroke Prevention in Atrial Fibrillation [SPAF-III] study). *J Am Soc Echocardiogr*. 1999;12(12):1080-7

⁸⁷ Takahashi N, Nakamura Y, Komatsu S, Kusano KF, Ohe T. The feasibility of tissue Doppler acceleration as a new predictor of thrombogenesis in the left atrial appendage associated with nonvalvular atrial fibrillation. *Echocardiography*. 2008;25(1):64-71

⁸⁸ Parvathaneni L, Mahenthiran J, Jacob S, et al. Comparison of tissue Doppler dynamics to Doppler flow in evaluating left atrial appendage function by transesophageal echocardiography. *Am J Cardiol*. 2005;95(8):1011-4

-
- ⁸⁹ Sallach JA, Puwanant S, Drinko JK, et al. Comprehensive left atrial appendage optimization of thrombus using surface echocardiography: the CLOTS multicenter pilot trial. *J Am Soc Echocardiogr.* 2009;22(10):1165-72
- ⁹⁰ Tamura H, Watanabe T, Hirono O, et al. Low wall velocity of left atrial appendage measured by trans-thoracic echocardiography predicts thrombus formation caused by atrial appendage dysfunction. *J Am Soc Echocardiogr.* 2010;23(5):545-552
- ⁹¹ Yoshida N, Okamoto M, Hirao H, et al. Role of transthoracic left atrial appendage wall motion velocity in patients with persistent atrial fibrillation and a low CHADS2 score. *J Cardiol* 2012;60(4):310-5
- ⁹² Tamura H, Watanabe T, Nishiyama S, et al. Prognostic value of low left atrial appendage wall velocity in patients with ischemic stroke and atrial fibrillation. *J Am Soc Echocardiogr.* 2012;25(5):576-83
- ⁹³ Masuda M, Iwakura K, Inoue K, et al. Estimation of left atrial blood stasis using diastolic late mitral annular velocity. *Eur Heart J Cardiovasc Imaging.* 2013;14(8):752-7
- ⁹⁴ Providência R, Faustino A, Ferreira MJ, et al. Evaluation of left atrial deformation to predict left atrial stasis in patients with non-valvular atrial fibrillation - a pilot-study. *Cardiovasc Ultrasound.* 2013;11:44
- ⁹⁵ Muellerleile K, Sultan A, Groth M, et al. Velocity encoded cardiovascular magnetic resonance to assess left atrial appendage emptying. *J Cardiovasc Magn Reson.* 2012;14:39

FIGURE LEGENDS

Figure 1. Late-gadolinium enhanced (LGE) magnetic resonance images representing fibrotic areas of left atrium as seen in antero-posterior view. The four images show different stages of increasing LGE burden (Utah classes, divided by quartiles): stage I (**A**: 2.7% enhancement), Stage II (**B**: 10.2% enhancement), Stage III (**C**: 19.4%), and Stage IV (**D**: 38.4% enhancement). Left atrial enhancement is represented as **green** areas to enhance contrast with non-enhanced tissue. Higher stages have been associated with LA appendage thrombi and history of previous stroke (see text for details). Reproduced with permission from Daccarett M et al. J Am Coll Cardiol 2011;57:831-8.

Figure 2. Left atrial appendage morphologies as represented in order of increasing complexity, which has been reported to relate to a growing burden of silent cerebral ischemia: from the simpler, Cactus (**A**, a dominant central lobe with small chambers extending in all directions), to the more complex, Cauliflower (**D**, complex internal characteristics with lack of a dominant lobe). Intermediate forms Chicken Wing (**B**, an obvious bend in the proximal or middle part of the dominant lobe) and Wind Sock (**C**, a dominant lobe plus secondary or even tertiary lobes arising from the dominant lobe) are as well represented. Reproduced with permission from Anselmino M et al. Heart Rhythm 2014;11(1):2-7.

Figure 3. Left atrial appendage wall velocity (TTE-LAWV) as measured by trans-thoracic tissue Doppler imagin (DTI). Placing sample volume at left atrial appendage (LAA) tip from the parasternal short-axis view in diastole, DTI velocities were obtained and the peak wall velocity of downward atrial waveform within each RR interval was averaged. Reduced TTE-LAWV related to history of previous stroke and predicted recurrent cerebrovascular events at follow up in those same patients⁹². AoV, Aortic valve. Reproduced with permission from Tamura H et al. J Am Soc

Figure 1.

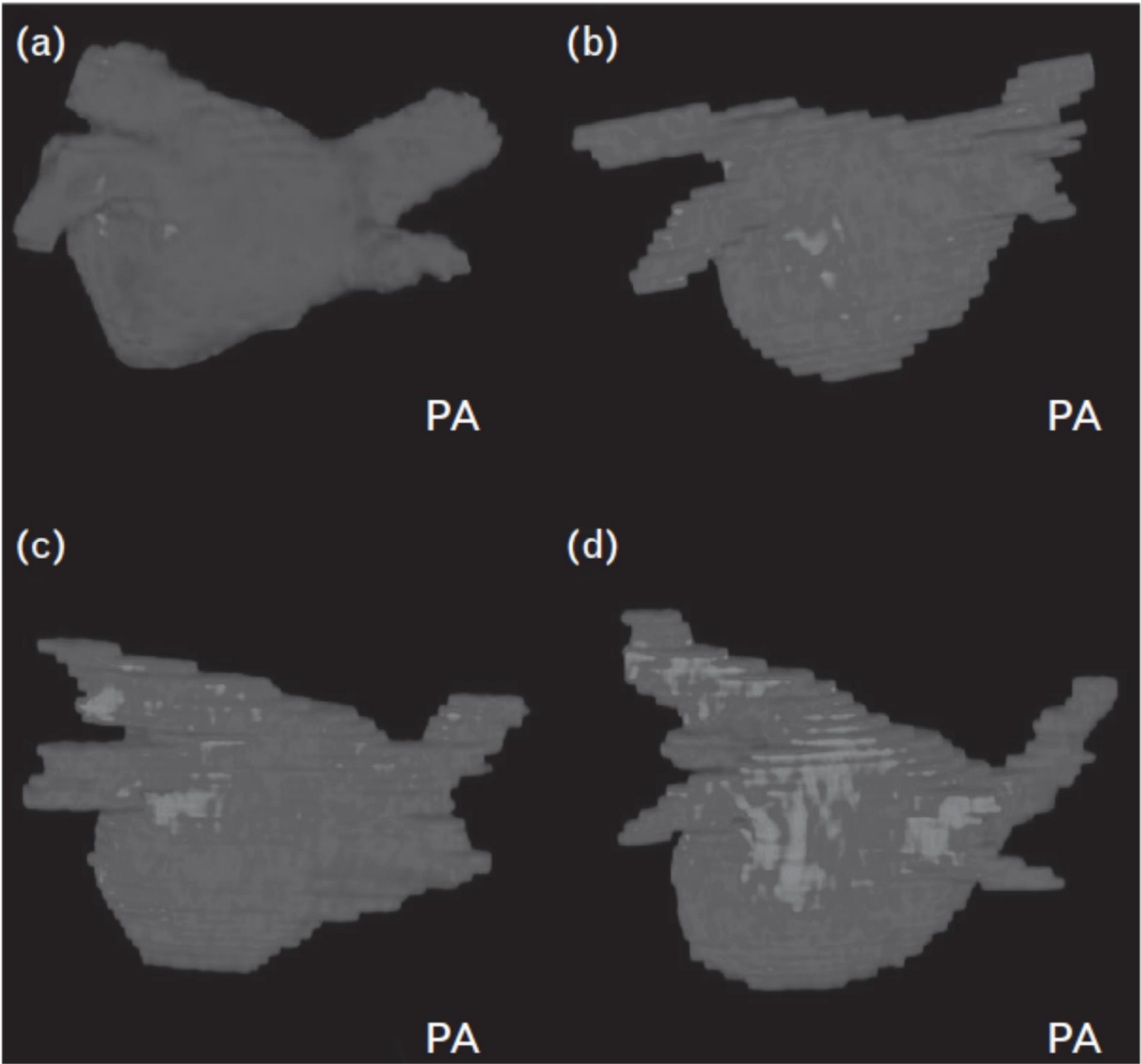


Figure 2.

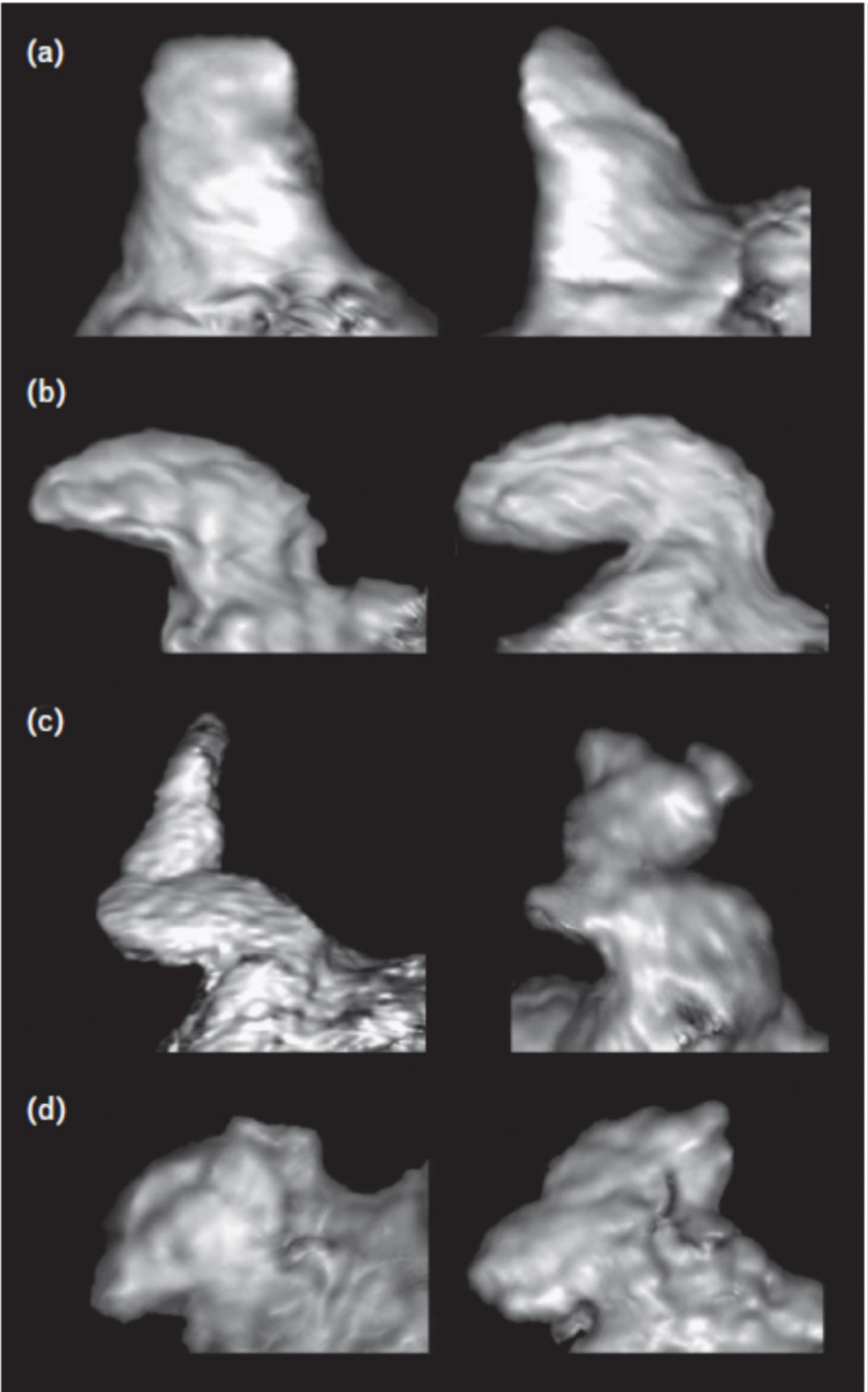


Figure 3.

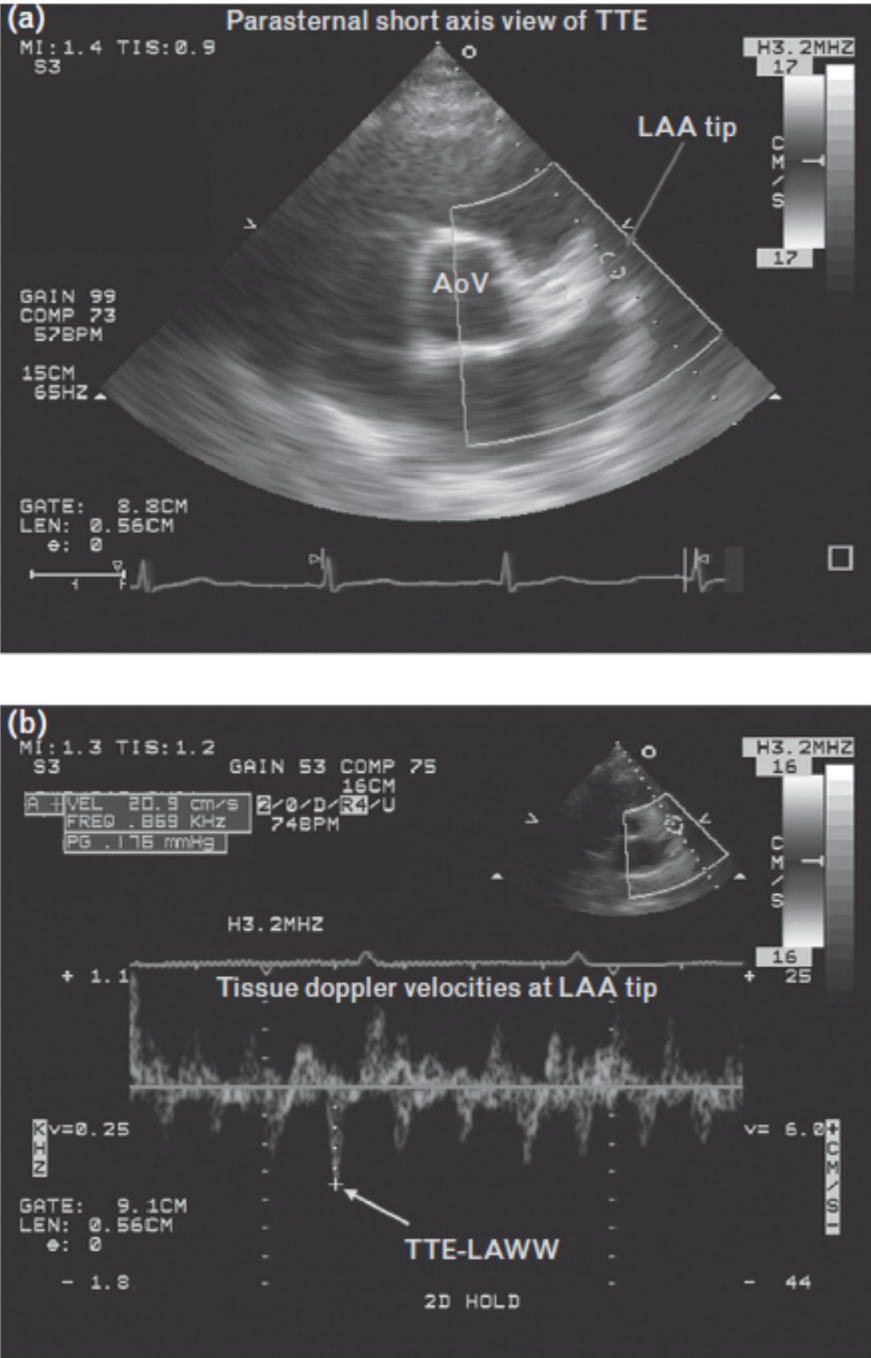


Table 1. Left atrial and appendage parameters tested for thromboembolic risk prediction in AF patients.

Parameter	No. of patients	Strength	Reproducibility	End-point	Feasibility	Technical limitations	Stroke prediction discordance
Left atrium AP diameter	+++	+	+++	+++	+++		Documented for nonatrial fibrillation patients; controversial for atrial fibrillation patients
Left atrium volume	+++	++	++	+++	+++		Documented for nonatrial fibrillation patients; controversial for atrial fibrillation patients
LAA size	–	+/-	–	+	+/-	Wide variability of methods and results	
LAA morphology	+	+	+/-	+	+/-	Low reproducibility of morphological stratification; MRI or angio-CT required (except one study performed by three-dimensional TEE)	
Left atrium LGE	+	+++	?	+	+/-	Lack of standardized algorithm; gadolinium enhanced-MRI required	One single study
Left atrium strain	+	+++	++	+	+++	Speckle tracking technique required	Only surrogate end points
LAAv	++	+++	+++	+++	+	TEE required	
LAA strain	+/-	++	++	++	+++	Not computable in 100% of the cases (TTE with TDI in >90% of cases)	Limited data available